Please amend the claims as follows. Deleted matter is indicated with strikethroughs and bolded text and new text is indicated with bolded and underlined text. These claims supersede all previous versions. No new matter is added by any of the changes.

IN THE CLAIMS:

1. (Currently Amended) A medical article An implantable or insertable medical device

comprising a release region, said release region comprising (a) a polymeric carrier comprising a

first polymer and (b) drug loaded nanoparticles dispersed within said polymeric carrier, said drug

loaded nanoparticles comprising: silicate particles comprising a layered silicate material; and a

first therapeutic agent, wherein the first therapeutic agent is structurally associated with the

silicate particles in that the first therapeutic agent occupies spaces between adjacent layers of the

silicate material of each silicate particle to form a depot for the first therapeutic agent.

2. (Currently Amended) The medical article device of claim 1, wherein said first therapeutic

agent is a hydrophilic therapeutic agent and said first polymer is a hydrophobic polymer.

3. (Currently Amended) The medical article device of claim 2, wherein said medical article

device is a vascular medical device, wherein said first therapeutic agent is halofuginone HBr, and

wherein said first polymer is a polyolefin-polyvinylaromatic block copolymer.

4. (Withdrawn and Currently Amended) The medical article device of claim 1, wherein said first

therapeutic agent is a hydrophobic therapeutic agent and said first polymer is a hydrophilic

polymer.

5. (Withdrawn and Currently Amended) The medical article device of claim 1, further comprising

a second polymer.

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6. (Withdrawn and Currently Amended) The medical article device of claim 5, wherein said

polymeric carrier further comprises said second polymer.

7. (Withdrawn and Currently Amended) The medical article device of claim 5, wherein said

nanoparticles further comprise said second polymer.

8. (Withdrawn and Amended) The medical article device of claim 7, wherein said second polymer

is hydrophobic and said first polymer is hydrophilic.

9. (Withdrawn and Currently Amended) The medical article device of claim 7, wherein said

second polymer is hydrophilic and said first polymer is hydrophobic.

10. (Withdrawn and Currently Amended) The medical article device of claim 9, wherein said

medical article is a vascular medical device, wherein said first therapeutic agent is

halofuginone HBr, wherein said first polymer is a polyolefin-polyvinylaromatic block copolymer,

and wherein said second polymer is a hydrophilic polymer selected from hyaluronic acid,

collagen, heparin, chrondroitin sulfate, phosphoro choline, dextran, and polyethylene oxide.

11. (Withdrawn and Currently Amended) The medical article device of claim 1, wherein said

polymeric carrier further comprises said first therapeutic agent.

12. (Withdrawn and Currently Amended) The medical article device of claim 1, further

comprising a second therapeutic agent.

13. (Withdrawn and Currently Amended) The medical article device of claim 12, wherein said

polymeric carrier further comprises said second therapeutic agent.

14. (Withdrawn and Currently Amended) The medical article device of claim 13, wherein said

first therapeutic agent is hydrophilic and said second therapeutic agents is hydrophobic.

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15. (Withdrawn and Currently Amended) The medical article device of claim 12, wherein said

nanoparticles further comprise said second therapeutic agent.

16. (Withdrawn and Currently Amended) The medical article device of claim 15, wherein said

first and second therapeutic agents are hydrophilic.

17. (Withdrawn and Currently Amended) The medical article device of claim 1, wherein said

release region is disposed over at least a portion of a medical article substrate.

18. Cancelled.

19. (Currently Amended) The medical article device of claim 18, wherein said implantable or

insertable medical device is adapted for implantation or insertion into the coronary or peripheral

vasculature.

20. (Withdrawn and Currently Amended) The medical article device of claim 19, wherein said

implantable or insertable medical device is adapted for implantation or insertion into the

esophagus, trachea, colon, biliary tract, urinary tract, prostate or brain.

21. (Currently Amended) The medical article device of claim 19, wherein said implantable or

insertable medical device is selected from a catheter, a guide wire, a balloon, a filter, a stent, a

stent graft, a vascular graft, a vascular patch, a shunt, an electrode, a heart valve, a circulation

pump, and an intraluminal paving system.

22. (Currently Amended) The medical article device of claim 19, wherein said therapeutic agent

is selected from an anti-thrombotic agent, an anti-proliferative agent, an anti-inflammatory agent,

an anti-migratory agent, an agent affecting extracellular matrix production and organization, an

antineoplastic agent, an anti-mitotic agent, an anesthetic agent, an anti-coagulant, a vascular cell

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growth promoter, a vascular cell growth inhibitor, a cholesterol-lowering agent, a vasodilating

agent, and an agent that interferes with endogenous vasoactive mechanisms.

23. (Currently Amended) The medical article device of claim 1, wherein said layered silicate

material comprises synthetic or naturally occurring smectite.

24. (Withdrawn and Currently Amended) The medical article device of claim 1, wherein said

layered silicate material comprises a natural or synthetic silicate material selected from bentonite,

aliettite, vermiculite, swinefordite, montmorillonite, yakhontovite, nontronite, beidellite,

volkonskoite, stevensite, hectorite, saponite, laponite, sauconite, magadiite, kenyaite and ledikite.

25. (Currently Amended) A method of releasing a therapeutic agent to a patient comprising: (a)

providing the medical article device of claim 1; and (b) contacting said medical article with a

patient.

26. (Withdrawn and Currently Amended) A method of providing the medical article device of

claim 1 comprising:

providing a release-region-forming fluid comprising (a) said first polymer species and (b)

said drug loaded nanoparticles; and

applying said release-region-forming fluid to a medical article substrate or to a releasable

template.

New Claims:

Please add new Claims 27 and 28 as follows:

27. (New) The medical device of claim 1 wherein the silicate particles have a maximum cross-

sectional length between 30 to 500 nm and spacing between the adjacent layers within the silicate

particles is in the range of 5-20Å.

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28. (New) The medical device of claim 2 wherein said hydrophobic polymer is selected from the group consisting of olefin polymers and copolymers, styrene polymers and copolymers, halogenated hydrocarbon polymers and copolymers, vinyl polymers and copolymers, polymers and copolymers of acrylic acid esters, polymers and copolymers of methacrylic acid esters, polycarbonates, polyimides, polyetheretherkeones, polyamides, polyvinylaceteates, polysulfones, polyethersulfones, polyethersulfones, polyethersulfones, polyurethanes and siloxane-urethane copolymers, and polyorganosiloxanes.